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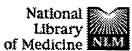
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Alternative conformations of amyloidogenic proteins govern their behavior.

Kelly JW.

Department of Chemistry, Texas A&M University, College Station 77843-3255, USA. kelly@chemvx.tamu.edu

Recent publications strongly support the hypothesis that conformational changes in amyloidogenic proteins lead to amyloid fibril formation and cause disease. Biophysical studies on several amyloidogenic proteins provide insights into the conformational changes required for fibrilogenesis. In addition, newly available moderate to high resolution structural studies are bringing us closer to understanding the structure of amyloid.

**Publication Types:** 

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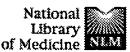
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Amyloid fibril formation and protein misassembly: a structural quest for insights into amyloid and prion diseases.

Kelly JW.

Department of Chemistry, Texas A&M University, College Station, Texas, 77843-3255, USA. kelly@chemvx.tamu.edu

The assembly and misassembly of normally soluble proteins into fibrilar structures is thought to be a causative agent in a variety of human amyloid and prion diseases. Structural and mechanistic studies of this process are beginning to elucidate the conformational changes required for the conversion of a normally soluble and functional protein into a defined quaternary structure.

Publication Types:

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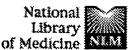
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## Amyloidosis.

## Tan SY, Pepys MB.

Department of Medicine, Royal Postgraduate Medical School, Hammersmith Hospital, London, UK.

Amyloidosis is a heterogeneous group of disorders characterized by extracellular deposition of abnormal protein fibrils which are derived from different proteins in different forms of the disease. Asymptomatic amyloid deposition in a variety of tissues is a universal accompaniment of ageing, and clinical amyloidosis is not rare. Intracerebral and cerebrovascular beta-protein amyloid deposits are a hallmark of the pathology of both sporadic and familial Alzheimer's disease, beta 2-microglobulin-derived amyloid is a common complication of long term haemodialysis, and islet amyloid polypeptide is the fibril protein in the universal islet amyloidosis of type II diabetes mellitus. New fibril proteins have lately been identified in hereditary amyloidosis, including variants of gelsolin, apolipoprotein AI, lysozyme and fibrinogen. The development of radiolabelled serum amyloid P component (SAP) scintigraphy has allowed amyloid to be diagnosed non-invasively in vivo for the first time, provided unique insight into the distribution and size of amyloid deposits, and yielded novel information on the natural history and the effects of treatment. Amyloid deposits are in a state of dynamic turnover and can regress if new fibril formation is halted. The recent elucidation of the three dimensional structure of human SAP may enable the design of specific therapeutic agents.

## Publication Types:

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Powerful solvent systems useful for synthesis of sparingly-soluble peptides in solution.

Kuroda H, Chen YN, Kimura T, Sakakibara S.

Peptide Institute Inc., Protein Research Foundation, Osaka, Japan.

Our maximum protection strategy for the synthesis of human parathyroid hormone(1-84) indicates that fully protected peptide segments in the form of Boc-peptide phenacyl (Pac) ester are relatively soluble in ordinary organic solvents such as DMF, NMP or DMSO, which are suitable for coupling segments. However, about 1% of such segments synthesized were found to be insoluble even in the most polar solvent, DMSO. Thus, a more powerful solvent which can be used for their peptide synthesis was pursued. Among the solvent systems tested, a mixture of trifluoroethanol (TFE) or hexafluoroisopropanol (HFIP) and trichloromethane (TCM) or dichloromethane (DCM) was found to be most powerful for dissolving such sparingly-soluble protected peptides. These solvent systems were confirmed to be useful for the removal reaction of the carboxy-terminal Pac esters from the sparingly-soluble segments. They were then tested for the coupling reactions of fully protected Boc-peptides with other sparingly-soluble peptide esters. The TFE/TCM or TFE/DCM system was extremely useful for coupling segments without danger of racemization and of trifluoroester formation, if WSCI was used as the coupling reagent in the presence of 3,4-dihydro-3hydroxy-4-oxo-1,2,3-benzotriazine (HOOBt).

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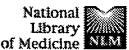
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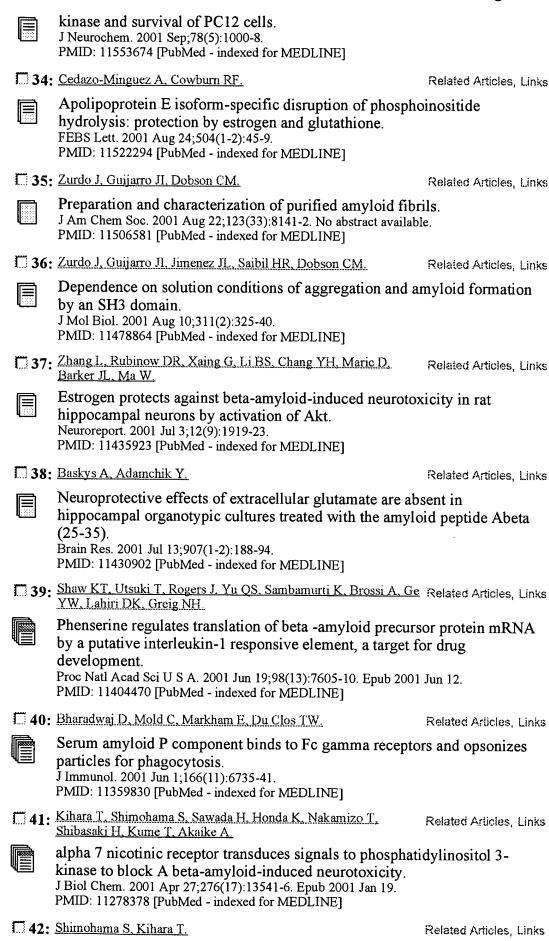
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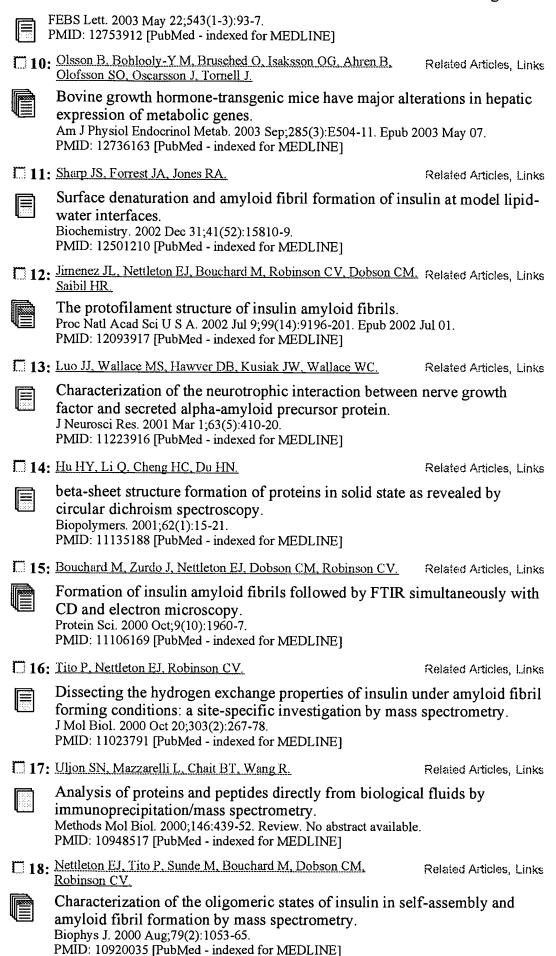
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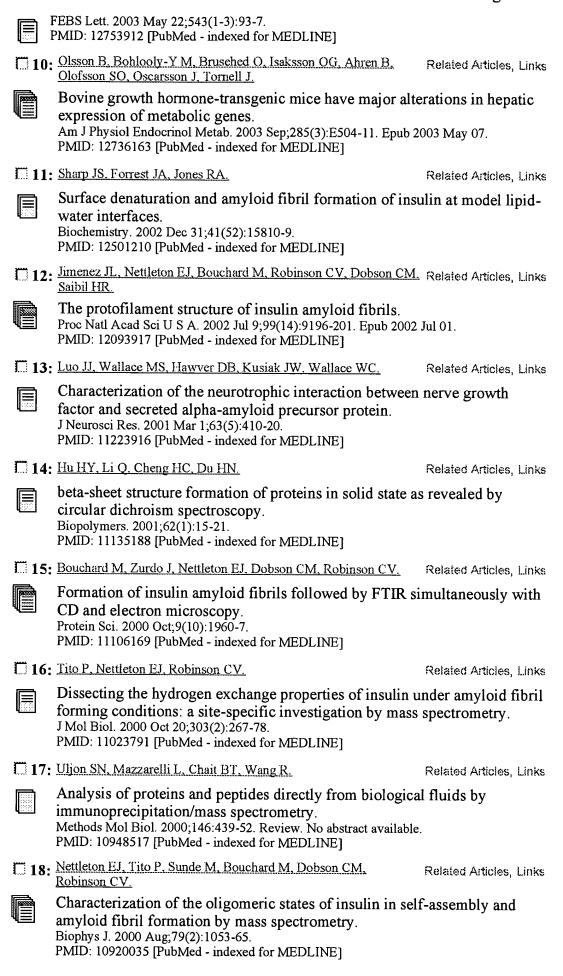
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# Helium Balloon

Can we use the <u>Ideal Gas Law</u>, PV=nRT, to calculate the lifting potential of a helium balloon?

- If two balloons are filled to equal volumes, the number of moles of gas molecules in one balloon will equal the moles of gas molecules in the other, even if the balloons are filled with different types of gas.
- If the weight per unit volume of a gas is less than the weight per unit volume of air, than the balloon will rise, provided that the difference in weights exceeds the weight of the balloon.
- Assume air to be 80% nitrogen and 20% oxygen.

With any problem, we have to ask ourselves what variables are readily available.

- temperature can be read from a thermometer
- pressure can be read from a barometer

What variables do we have to experimentally determine? What variables can be calculated?

Weigh the balloon, the ropes, and the basket.

Assume that today the weather gives us STP conditions: standard temperature (25 deg C) and standard pressure (1 atm.) To keep things simple, assume that these variables don't change.

Later, we may wish to calculate the effect (if any) of temperature on lift power.

Convert temperature from Celcius to Kelvin since we need an absolute temperature scale for calculations involving gas laws.

$$25 C + 273.15 = 298.15 K$$

Since temperature has the units of Kelvin and pressure has the units of atmospheres, we will want to use the real gas constant

$$R = 0.08206 L atm K^{-1} mol^{-1}$$

Note: you may find real gas constants with different numbers, other than 0.08206. If so, notice that the units are different. Using <u>dimensional analysis</u> it is possible to change units, and thus convert from one real gas constant to another.

After we have determined V, it is possible to solve the Ideal Gas Law for moles, n.

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We assume pressure and temperature are constant (<u>More information about assumptions</u>). We have determined volume using a method not specified. We know that R is a constant. Since all variables are set, there is only one possible answer for moles, n.

If we fill the balloon with any gas to the above specifications (T = 25 C and P = 1 atm), once full, there will be the same number of moles of the gas in the balloon, regardless of what type of gas the ballon is filled with.

If the weight of the balloon filled with a gas is less than the weight of the balloon filled with 20% oxygen and 80% nitrogen, then the difference in mass will correspond to a bouyant force.

If the difference in weight between the balloon filled with air, and the balloon filled with a lighter gas is greater than the sum of the weight of the balloon, the ropes, the basket, and your weight, then the balloon will carry you.

#### Example:

- o A large balloon is filled with 1000 lbs of air.
- o When filled with helium, the helium in the balloon weighs 200 lbs.
- o The balloon is capable of lifting 800 lbs.
- o The balloon weights 300 lbs.
- o The ropes weigh 50 lbs.
- o The basket weighs 200 lbs.
- o 300 lbs + 50 lbs + 200 lbs = 550 lbs.
- o 800 lbs 550 lbs = 250 lbs.
- o The balloon is capable of lifting 250 lbs.

Assume that the balloon, the rope, and the basket have a mass of 30 kilograms, and your mass is 70 kilograms. Use helium for the gas. Hydrogen is lighter, but recall the Hindenburg disaster. Assume that the volume of the balloon is 1000 L.

Mass of displaced air:

Mass of 800 L of nitrogen:

PV 1.0000 atm \* 800.00 L   

$$n = -- = ----- = 32.698$$
 moles   
RT 0.08206 (L atm)/(mol K) \* 298.15 K

Mass of 200 L of oxygen:

15.9994 g 
$$\rm O_2$$
 8.1745 moles  $\rm O_2$  \* ------ = 130.79 g  $\rm O_2$  mole  $\rm O_2$ 

Mass of helium:

1000 L of helium:

Mass of air: 457.99 g + 130.79 q = 588.78 g

Mass of air - Mass of helium = 588.78 g - 163.6 g = 425.2 g

425.2 g isn't much, roughly equivalent to a pound (1 lb = 453.59 g)

As a homework problem, confirm that 1000 L corresponds to a cubic meter.

Thought question: "What might go wrong if the balloon is too big (where 'too big; signifies "big enough to create the problem hypothesized).

#### More information about the assumptions

We assume that there is no leak of helium out of the balloon, or nitrogen or oxygen into the balloon. If helium could leak out, then over time the management of the balloon of the balloon.

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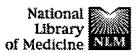
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In vitro synthesis of "amyloid" fibrils from insulin, calcitonin and parathormone.

Kedar I, Ravid M, Sohar E.

Insulin, calcitonin and parathyroid hormone subjected to one of two procedures-acidification and heating or incubation with mouse kidney lysosomal extracts-assumed a nonbranching fibrillar structure, 7 to 10 nm in diameter. The preparations showed green birefringence after Congo red staining. The in vitro synthesis from different hormonal polypeptides of fibrils, fulfilling the criteria for the identification of amyloid, indicates that these criteria are related to conformational rather than to compositional properties, and suggests that these hormones may provide the subunit of the amyloid formed in the corresponding endocrine organs.

PMID: 62581 [PubMed - indexed for MEDLINE]

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# Amyloid in polypeptide hormone-producing tumors.

Westermark P, Grimelius L, Polak JM, Larsson LI, Van Noorden S, Wilander E, Pearse AG.

The hormone content of 72 endocrine tumors was determined by immunofluorescence and their amyloid content was investigated. Seventeen of the 72 tumors contained amyloid. Amyloid was frequently found in tumors producing calcitonin, insulin, or growth hormone, but was rarely found in other tumors. Thus, there is a relationship between the occurrence of amyloid in an endocrine tumor and the type of hormone it produces. The reason for this is not known, but there is evidence that the amyloid fibrils contain proteins related to the hormone produced by the tumors.

PMID: 881783 [PubMed - indexed for MEDLINE]

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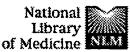
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# Iatrogenic, insulin-dependent, local amyloidosis.

### Storkel S, Schneider HM, Muntefering H, Kashiwagi S.

Human and experimental amyloidosis can occur either as a generalized widespread deposit of various proteins or a localized deposit. We looked for local amyloidosis caused iatrogenically under clinical and experimental conditions. Subcutaneous tissue from one diabetic patient and six Wistar rats, which had received a continuous local infusion of 1.2 iu of insulin daily for 6 weeks, was examined histologically. In all cases the development of granulation tissue around the tip of the catheter was observed. In addition, inhomogenous extracellular deposits showing green birefringence under polarized light when stained Congo red were seen. Immunohistologically, they displayed binding of anti-insulin antibody. Electron microscopy demonstrated a typical spear-like fibrillar structure with a fibril diameter of 60 to 80 A. These findings confirmed that the deposited substance was amyloid. Iatrogenically administered protein produced in vivo amyloidosis at the site of its entry. Insulin can lead to the formation of amyloid fibrils not only in vitro but also in vivo.

PMID: 6337294 [PubMed - indexed for MEDLINE]

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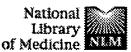
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# [Amyloidosis of the pancreatic islets and diabetes mellitus]

[Article in Russian]

### Ageev AK.

Pancreas was examined in 136 patients who died at the age of 7 to 89 years of various diseases including 22 with diabetes mellitus. Amyloidosis of its islands was observed in 9 patients (aged 49 and over); 6 out of them suffered from diabetes mellitus. Number of islands with amyloidosis and amyloid quantity were determined morphometrically. Glucagon-producing A-cells and insulin-producing B-cells in the islands not involved in amyloidosis were counted in sections impregnated by Grimelius. It is found that the development of diabetes is determined not only by the islands amyloidosis but by the quantitative domination of A-cells over B-cells in the islands without amyloidosis as well being the manifestation of aging processes.

PMID: 3527115 [PubMed - indexed for MEDLINE]

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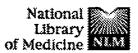
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Quantitative evaluation of congo red binding to amyloid-like proteins with a beta-pleated sheet conformation.

Klunk WE, Pettegrew JW, Abraham DJ.

Department of Psychiatry, University of Pittsburgh School of Medicine, Pennsylvania.

The binding of Congo red to several purified amyloid-like peptides having a beta-pleated sheet conformation was quantitatively examined. Congo red binds preferentially to the beta-pleated sheet conformation of both insulin fibrils and poly-L-lysine. Congo red does not bind nearly so well to poly-Lserine or polyglycine, despite the fact that these peptides also have a betapleated sheet conformation. Binding to insulin fibrils was saturable with an apparent Bmax of 2 moles of Congo red per mole of insulin fibrils and an apparent KD of 1.75 x 10(-7) M. Binding to beta-poly-L-lysine was similar but had a much higher apparent Bmax of 43. Binding of Congo red to betapoly-L-lysine was pH dependent and appeared to be determined by the number of protonated lysine residues in the 250 amino acid peptide. We present a new hypothesis in which Congo red binds to amyloid-like proteins via bonds between the two negatively charged sulfonic acid groups of Congo red and two positively charged amino acid residues of two separate protein molecules which are properly oriented by virtue of the beta-pleated sheet conformation of the peptide backbone.

PMID: 2666510 [PubMed - indexed for MEDLINE]

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Binding of the dye congo red to the amyloid protein pig insulin reveals a novel homology amongst amyloid-forming peptide sequences.

Turnell WG, Finch JT.

MRC Laboratory of Molecular Biology, Cambridge, U.K.

The three-dimensional structure has been determined of a complex of the dye Congo Red, a specific stain for amyloid deposits, bound to the amyloid protein insulin. One dye molecule intercalates between two globular insulin molecules at an interface formed by a pair of anti-parallel beta-strands. This result, together with analysis of the primary sequences of other amyloidogenic proteins and peptides suggests that this mode of dye-binding to amyloid could be general. Moreover, the structure of this dye-binding interface between protein molecules provides an insight into the polymerization of amyloidogenic proteins into amyloid fibres. Thus the detailed characterization, at a resolution of 2.5 A, of the dye binding site in insulin could form a basis for the design of agents targeted against a variety of amyloid deposits.

PMID: 1433294 [PubMed - indexed for MEDLINE]

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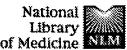
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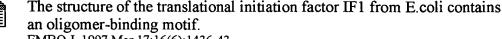
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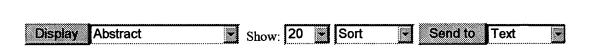
Overproduction, crystallization, and preliminary X-ray diffraction studies of the major cold shock protein from Bacillus subtilis, CspB.

Schindelin H, Herrler M, Willimsky G, Marahiel MA, Heinemann U.

Institut für Kristallographie, Freie Universität Berlin, Federal Republic of Germany.

The major cold shock protein from Bacillus subtilis (CspB) was overexpressed using the bacteriophage T7 RNA polymerase/promoter system and purified to apparent homogeneity from recombinant Escherichia coli cells. CspB was crystallized in two different forms using vapor diffusion methods. The first crystal form obtained with ammonium sulfate as precipitant belongs to the trigonal crystal system, space group P3(1)21 (P3(2)21) with unit cell dimensions a = b = 59.1 A and c = 46.4 A. The second crystal form is tetragonal, space group P4(1)2(1)2 (P4(3)2(1)2) with unit cell dimensions a = b = 56.9 A and c = 53.0 A. These crystals grow with polyethylene glycol 4000 as precipitant.

PMID: 1409560 [PubMed - indexed for MEDLINE]



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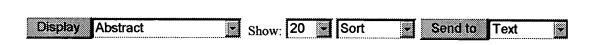
Effect of pH and phosphate ions on self-association properties of the major cold-shock protein from Bacillus subtilis.

Makhatadze GI, Marahiel MA.

Department of Biology, Johns Hopkins University, Baltimore, Maryland 21218

The intermolecular interactions of the major cold-shock protein from Bacillus subtilis (CspB) in solution in the presence of different salts, including phosphate, have been studied by means of scanning calorimetry and sizeexclusion chromatography. Calorimetric results indicate that, in all cases, protein unfolding can be approximated by a 2-state model, but the modes of unfolding can differ depending on the conditions. In the presence of phosphate, the cooperative folding unit is a monomer, whereas in the absence of phosphate, the cooperative unit is a dimer. The difference in the selfassociation of CspB in the presence and absence of phosphate was supported by size-exclusion chromatography. These results are compared with recent structural studies of CspB in crystal and in solution.

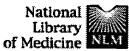
PMID: 7703860 [PubMed - indexed for MEDLINE]



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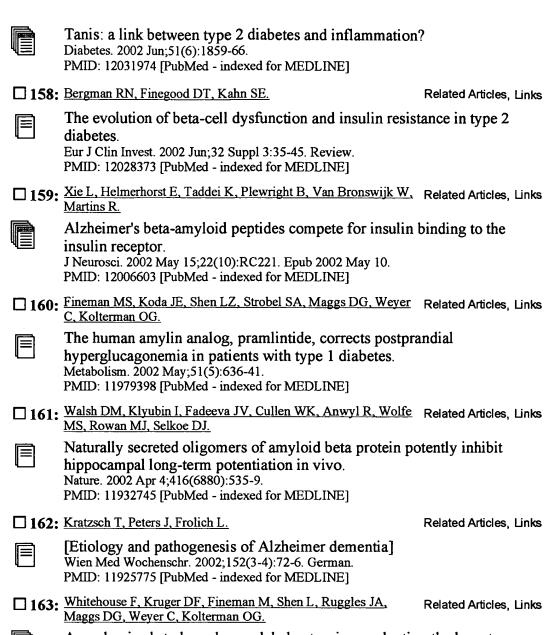
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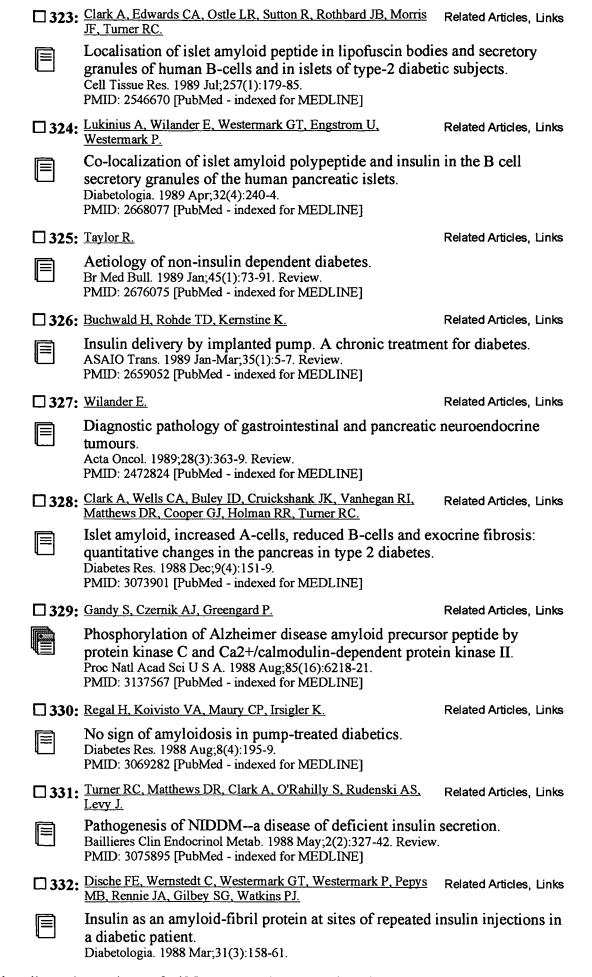
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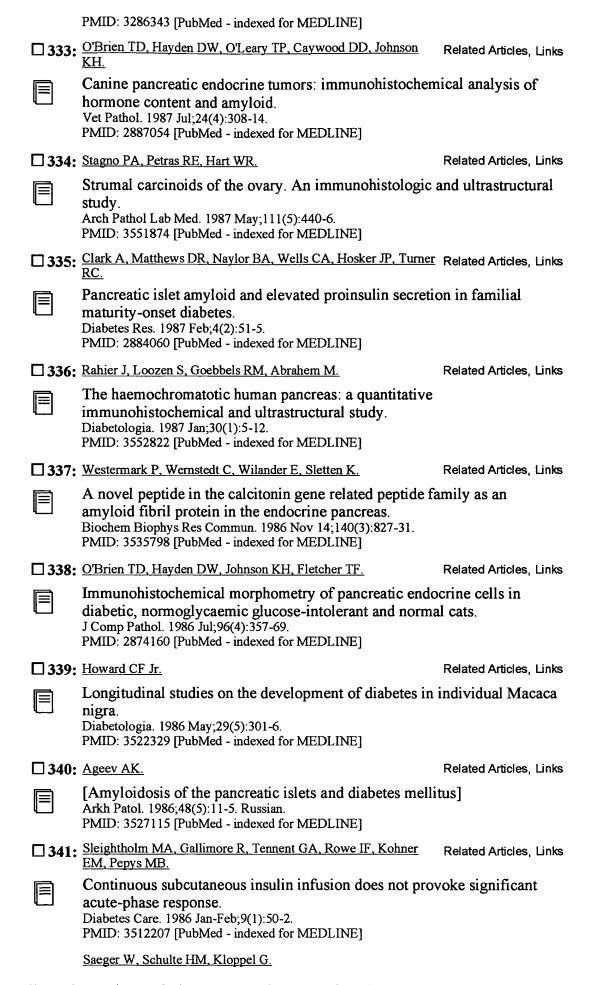
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1996:462550 CAPLUS
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      Antibodies and antibody fragments for prevention of protein aggregation
TI
      and therapy of diseases associated therewith
      Solomon, Beka
IN
      Ramot-Univ. Authority for Applied Research and Industrial Development
PA
      Ltd., Israel; Shoshan, Herbert Z. PCT Int. Appl., 60 pp.
SO
      CODEN: PIXXD2
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RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG
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L11
      1994:264491 CAPLUS
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DN
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      Prediction of the active sites of proteins from amino acid sequences
TI
      Numao, Naganori; Kidokoro, Shunichi
Sagami Chem. Res. Cent., Sagamihara, 229, Japan
Biological & Pharmaceutical Bulletin ( ***1993*** ), 16(11), 1160-3
AU
CS
SO
      CODEN: BPBLEO; ISSN: 0918-6158
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      Journal
LΑ
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L11
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      1993:76636
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      118:76636
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      Method for surmising functional site in physiologically active polypeptide
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      or polynucleotide
      Numao, Naganori; Kidokoro, Shunichi
Sagami Chemical Research Center, Japan; Tosoh Corp.; Nippon Mining Co.,
IN
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      Eur. Pat. Appl., 75 pp.
      CODEN: EPXXDW
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IN
        Tunq J S
                       ATHENA NEUROSCIENCES INC.
PA
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                       CHRYSLER S.
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                       MCCONLOGUE L.
        (SINH-I)
                       SINHA S.
                       TATSUNO G.
         (TATS-I)
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        (TUNG-I)
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TI
        beta- ***amyloid*** peptide prodn. in cells, use in Alzheimer's disease, also prepn. of cathepsin Y and nucleic acid encoding for it. Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
IN
                        ATHENA NEUROSCIENCES INC.
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PA
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TI
IN
        Tung J S
         (ATHE-N)
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                        ATHENA NEUROSCIENCES INC.
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        Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
IN
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        Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
IN
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TI
        Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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TI
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IN
        Tung J S
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        (ANDE-I)
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       Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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       ANSWER 47 OF 125 DGENE AAW93378 Protein
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AN
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TI
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       Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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       Acylamino and acyl:peptido:amino alcohol and aldehyde derivs. - inhibit beta- ***amyloid*** peptide prodn. in cells, use in Alzheimer's
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IN
       Tung J S
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       Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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       Acylamino and acyl:peptido:amino alcohol and aldehyde derivs.
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       Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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        (SINH-I)
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ΤI
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        (ATHE-N)
PA
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                      ANDERSON J.
        (ANDE-I)
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IN
                      ATHENA NEUROSCIENCES INC.
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PA
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       Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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PA
        (ATHE-N)
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        Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
        Tung J S
                       ATHENA NEUROSCIENCES INC.
PA
        (ATHE-N)
        (ANDE-I)
                       ANDERSON J.
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        Anderson J; Chrysler S; McConlogue L; Semko C M F; Sinha S; Tatsuno G;
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          Fibrinogen-coated particles for therapeutic use
Yen, Richard C. K., Yorba Linda, CA, United States
Hemosphere, Inc., Anaheim, CA, United States (U.S. corporation)
TI
IN
PA
                                                20020521
PI
           US 6391343
                                        B1
                             19961212
           WO 9639128
           US 1998-952765
                                                19980410 (8)
AΙ
                                                19960604
           WO 1996-US9458
                                                19980410
                                                               PCT 371 date
          Continuation-in-part of Ser. No. US 1995-554919, filed on 9 Nov 1995, now abandoned Continuation-in-part of Ser. No. US 1995-471650, filed on 6 Jun 1995, now patented, Pat. No. US 5725804 Continuation-in-part of Ser. No. US 1994-212546, filed on 14 Mar 1994, now patented, Pat. No. US 5616311 Continuation-in-part of Ser. No. US 1993-69831, filed on 1 Jun 1993, now abandoned Continuation-in-part of Ser. No. US 1992-959560, filed on 13 Oct 1992, now patented, Pat. No. US 5308620 Continuation-in-part of Ser. No. US 1991-641720, filed on 15 Jan 1991, now abandoned
RLI
           now abandoned
DT
           Utility
FS
           GRANTED
LN.CNT 2407
           INCLM: 424/491.000
INCLS: 424/078.060; 427/002.140; 514/002.000; 514/834.000; 514/937.000;
INCL
                      514/951.000; 516/077.000
           NCLM:
                      424/491.000
NCL
                      424/078.060; 427/002.140; 514/002.000; 514/834.000; 514/937.000;
           NCLS:
                      514/951.000; 516/077.000
           [7]
IC
           ICM: A61K009-16
           ICS: A61K038-36; A61K038-38
EXF 264/4.3; 427/2.14; 427/2.21; 427/213.3; 427/213.33; 424/78.06; 424/491; 424/493; 514/2; 514/834; 514/937; 514/951; 514/965; 516/77 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        ANSWER 84 OF 125
                                    USPATFULL on STN
L11
           2001:116764 USPATFULL
ΑN
           Ataxia-telangiectasia gene and its genomic organization
TI
IN
           Shiloh, Yosef, Tel Aviv, Israel
           Ramot-University Authority for Applied Research and Industrial Development, Tel Aviv, Israel (non-U.S. corporation)
PA
                                                20010724
PI
           US 6265158
                                        B1
           WO 9636691
                              19961121
                                                                                                          <--
           US 1998-952014
                                                19980202 (8)
AΙ
                                                19960516
           WO 1996-US7025
                                                19980202
                                                                PCT 371 date
                                                               PCT 102(e) date
                                                19980202
           Continuation-in-part of Ser. No. US 1996-629001, filed on 8 Apr 1996,
RLI
           now patented, Pat. No. US 5858661 Continuation-in-part of Ser. No. US
           1995-441822, filed on 16 May 1995, now patented, Pat. No. US 5756288
DT
           Utility
FS
           GRANTED
LN.CNT
           3109
           INCLM: 435/006.000
INCL
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NCLM:
                   435/006.000
NCL
                   536/023.100; 536/024.300; 536/024.310
         NCLS:
          [7]
IC
          ICM: C12Q001-68
          ICS: C07H021-04
          435/6; 536/23.1; 536/24.3; 536/24.31
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
       ANSWER 85 OF 125 USPATFULL on STN
                          USPATFULL
          2001:107872
AN
         Delivery of gene products by intestinal cell expression German, Michael, San Francisco, CA, United States
TI
IN
         Goldfine, Ira D., Kentfield, CA, United States
Rothman, Stephen S., Berkeley, CA, United States
         The Regents of the University of California, Oakland, CA, United States
PA
          (U.S. corporation)
         US 6258789
                                    B1
                                           20010710
PI
         WO 9811779
                          19980326
                                                                                               <--
ΑI
         US 1999-254988
                                           19990611 (9)
         WO 1997-US16523
                                           19970918
                                                         PCT 371 date PCT 102(e) date
                                           19990611
                                           19990611
          Continuation-in-part of Ser. No. US 1996-717084, filed on 20 Sep 1996
RLI
DT
          Utility
FS
          GRANTED
         1591
LN.CNT
INCL
          INCLM: 514/044.000
          INCLS: 435/320.100; 435/455.000; 435/458.000
                   514/044.000
NCL
         NCLM:
                   435/320.100; 435/455.000; 435/458.000
         NCLS:
IC
          [7]
          ICM: A61K048-00
514/44; 424/93.2; 424/93.21; 435/320.1; 435/455; 435/458; 435/325;
435/69.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 86 OF 125 USPATFULL on STN
L11
                         USPATFULL
AN
          2001:48208
          Ataxia-telangiectasia gene
TI
          Shiloh, Yosef, Tel Aviv, Israel
IN
         Tagle, Danilo A., Gaithersburg, MD, United States
Collins, Francis, Rockville, MD, United States
The United States of America as represented by the Department of Health
and Human Services, Washington, DC, United States (U.S. government)
Ramot University Authority for Applied Research and Industrial Dev.,
Israel (non-U.S. corporation)
PA
PΙ
          US 6211336
                                    B\bar{1}
                                           20010403
                          19961121
          WO 9636695
                                                                                               <--
          US 1998-952127
                                            19980226 (8)
ΑI
          WO 1996-US7040
                                            19960516
                                                         PCT 371 date
PCT 102(e) date
                                            19980226
         19980226 PCT 102(e) date
Continuation-in-part of Ser. No. US 1995-508836, filed on 28 Jul 1995, now patented, Pat. No. US 5777093 Continuation-in-part of Ser. No. US 1995-493092, filed on 21 Jun 1995, now patented, Pat. No. US 5728807
RLI
          Continuation-in-part of Ser. No. US 1995-441822, filed on 16 May 1995,
          now patented, Pat. No. US 5756288
DT
          Utility
FS
          Granted
LN.CNT
          2279
          INCLM: 530/350.000
INCLS: 530/326.000
NCLM: 530/350.000
NCLS: 530/326.000
INCL
NCL
          [7]
IC
          ICM: C07K001-00
          ICS: C07K014-00; C07K017-00
          530/326; 530/350
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 87 OF 125
                               USPATFULL on STN
L11
          1999:132587 USPATFULL
AN
TΙ
          Tryptase inhibitor
          Fritz, Hans, Icking, Germany, Federal Republic of
IN
          Sommerhoff, Christian, Munich, Germany, Federal Republic of
```

```
UCP Gen-Pharma AG, Zurich, Switzerland (non-U.S. corporation)
         US 5972698
ΡI
                                          19991026
         WO 9503333
                          19950202
         US 1996-586676
                                          19960125 (8)
ΑI
         WO 1994-EP2445
                                          19940725
                                          19960125
                                                        PCT 371 date
                                          19960125
                                                        PCT 102(e) date
PRAI
         EP 1993-111930
                                     19930726
DT
         Utility
FS
         Granted
LN.CNT
         1988
         INCLM: 435/320.100
INCL
         INCLS: 435/069.200; 435/212.000; 514/012.000; 530/324.000; 536/023.500
                   435/320.100
NCL
         NCLM:
         NCLS:
                   435/069.200; 435/212.000; 514/012.000; 530/324.000; 536/023.500
IC
          [6]
         ICM: C07K014-815
         ICS: C12N015-11; A61K038-58
         435/219; 435/69.2; 435/172.3; 435/320.1; 435/325; 435/252.3; 435/254.11; 514/2; 514/826; 530/300; 530/324; 536/23.1; 536/23.5
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 88 OF 125 USPATFULL on STN
L11
         1998:162469
                          USPATFULL
AN
         A.beta. peptides that modulate .beta.-
                                                                ***amyloid***
                                                                                      aggregation
TI
         Findeis, Mark A., Cambridge, MA, United States
Benjamin, Howard, Lexington, MA, United States
IN
         Garnick, Marc B., Brookline, MA, United States
Gefter, Malcolm L., Lincoln, MA, United States
         Hundal, Arvind, Brighton, MA, United States
Kasman, Laura, Athens, GA, United States
         Musso, Gary, Hopkinton, MA, United States
Signer, Ethan R., Cambridge, MA, United States
Wakefield, James, Brookline, MA, United States
Reed, Michael, Marietta, GA, United States
Molineaux, Susan, Brookline, MA, United States
Kubasek, William, Belmont, MA, United States
         Chin, Joseph, Salem, MA, United States
         Lee, Jung-Ja, Wayland, MA, United States
Kelley, Michael, Arlington, MA, United States
         Praecis Pharmaceuticals, Inc., Cambridge, MA, United States (U.S.
PA
         corporation)
US 5854204
US 1996-612785
PΙ
                                          19981229
                                          19960314 (8)
AΙ
         Continuation-in-part of Ser. No. US 1995-404831, filed on 14 Mar 1995 And a continuation-in-part of Ser. No. US 1995-475579, filed on 7 Jun
RLI
         1995 And a continuation-in-part of Ser. No. US 1995-548998, filed on 27
         Oct 1995
DT
         Utility
         Granted
FS
         4304
LN.CNT
INCL
          INCLM: 514/002.000
                   514/012.000; 514/014.000; 530/324.000; 530/326.000
514/002.000
          INCLS:
NCL
         NCLM:
                   514/012.000; 514/014.000; 530/324.000; 530/326.000
         NCLS:
IC
          [6]
          ICM: C07K014-435
          ICS: C07K007-08
          514/14; 514/12; 514/2; 530/300; 530/324; 530/326; 930/10
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 89 OF 125
                               USPATFULL on STN
L11
          1998:162337 USPATFULL
ΑN
          Hexokinase inhibitors
TI
         Newgard, Christopher B., Dallas, TX, United States
Han, He-Ping, Arlington, TX, United States
Normington, Karl D., Dallas, TX, United States
IN
          Board of Regents, The University of texas System, Austin, TX, United
PA
          States (U.S. corporation)
                      Inc., Dallas, TX, United States (U.S. corporation)
          Betagene,
                                          19981229
          US 5854067
PΙ
          US 1996-588983
                                          19960119 (8)
ΑI
DT
          Utility
FS
          Granted
LN.CNT 5377
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INCLS: 425/004.000; 425/006.000; 425/091.100; 425/091.310; 425/183.000;
                 425/320.100; 425/325.000; 536/023.100; 536/024.310; 536/024.500
NCL
                 435/366.000
        NCLM:
                 435/004.000; 435/006.000; 435/091.100; 435/091.310; 435/183.000;
        NCLS:
                 435/320.100; 435/325.000; 536/023.100; 536/024.310; 536/024.500
IC
         [6]
         ICM: C12N015-85
ICS: C12N015-00; C12N015-63; C12Q001-68

EXF 435/325; 435/4; 435/6; 435/69.1; 435/320.1; 435/172.3; 424/94.1; 536/23.1; 536/24.5; 514/44; 576/24.31

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 90 OF 125
                            USPATFULL on STN
         1998:157315
                        USPATFULL
AN
        Cathepsin and methods and compositions for inhibition thereof
TI
         Tung, Jay S., Belmont, CA, United States
IN
         Sinha, Sukanto, San Francisco, CA, United States
        McConloque, Lisa, San Francisco, CA, United States
        Semko, Christopher M. F., Fremont, CA, United States
Athena Neurosciences, Inc., South San Francisco, CA, United States (U.S.
PA
        corporation)
        US 5849711
US 1995-469362
ΡI
                                      19981215
                                                                                    <--
ΑI
                                      19950606 (8)
        Utility
DT
FS
         Granted
LN.CNT
        2445
INCL
        INCLM: 514/019.000
                 514/693.000; 514/706.000; 514/715.000; 514/716.000; 514/721.000;
         INCLS:
                 514/724.000; 514/727.000
NCL
        NCLM:
                 514/019.000
                 514/693.000; 514/704.000; 514/715.000; 514/716.000; 514/721.000; 514/724.000; 514/727.000
        NCLS:
IC
         [6]
         ICM: A61K038-06
         ICS: A01N035-00; A01N033-18; A01N031-00
EXF
         514/19; 514/693; 514/704; 564/123
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 91 OF 125
                            USPATFULL on STN
L11
        1998:150698
                       USPATFULL
ΑN
TI
        Dioxetane compounds for the chemiluminescent detection of proteases,
        methods of use and kits therefore
        Bronstein, Irena, Newton, MA, United States
Edwards, Brooks, Cambridge, MA, United States
Martin, Christopher, Belmont, MA, United States
Sparks, Alison, North Andover, MA, United States
IN
         Voyta, John C., Sudbury, MA, United States
         Tropix, Inc., New Bedford, MA, United States (U.S. corporation)
PA
                                      19981201
PΙ
        US 5843681
AΙ
        US 1996-728990
                                      19961011 (8)
         Continuation of Ser. No. US 1995-385788, filed on 9 Feb 1995, now
RLI
        patented, Pat. No. US 5591591
Utility
DT
FS
         Granted
LN.CNT
        764
INCL
         INCLM: 435/007.400
         INCLS: 435/006.000; 530/330.000; 530/331.000; 530/807.000; 548/526.000;
                 549/264.000; 549/332.000
NCL
        NCLM:
                 435/007.400
                 435/006.000; 530/330.000; 530/331.000; 530/807.000; 548/526.000; 549/264.000; 549/332.000
        NCLS:
IC
         [6]
         ICM: G01N033-573
         ICS: C07D321-00
         435/7.4; 435/6; 549/332; 549/264; 530/331; 530/807; 530/330; 548/526
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                            USPATFULL on STN
      ANSWER 92 OF 125
L11
AN
         1998:143904 USPATFULL
TI
         Directed evolution of novel binding proteins
         Ladner, Robert Charles, Ijamsville, MD, United States
IN
        Gutterman, Sonia Kosow, Belmont, MA, United States
Roberts, Bruce Lindsay, Milford, MA, United States
Markland, William, Milford, MA, United States
         Ley, Arthur Charles, Newton, MA, United States
```

```
Corp., Cambridge, MA, United States (U.S. corporation)
PA
          Dyax,
          บร์ 5837500
                                           19981117
PΙ
          US 1995-415922
                                           19950403 (8)
AΙ
          Continuation of Ser. No. US 1993-9319, filed on 26 Jan 1993, now
RLI
         patented, Pat. No. US 5403484 which is a division of Ser. No. US 1991-664989, filed on 1 Mar 1991, now patented, Pat. No. US 5223409 which is a continuation-in-part of Ser. No. US 1990-487063, filed on 2 Mar 1990, now abandoned which is a continuation-in-part of Ser. No. US
          1988-240160, filed on 2 Sep 1988, now abandoned
DT
          Utility
FS
          Granted
LN.CNT 15973
          INCLM: 435/069.700
INCL
          INCLS: 435/172.300; 530/350.000; 530/412.000; 536/023.400
                    435/069.700
NCL
          NCLM:
                    435/091.100; 435/091.200; 435/471.000; 530/350.000; 530/412.000;
          NCLS:
                    536/023.400
IC
          [6]
          ICM: C12N015-62
          ICS: C07K019-00
          435/69.7; 435/172.3; 530/350; 530/412; 536/23.4
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 93 OF 125 USPAT
1998:91815 USPATFULL
                                USPATFULL on STN
L11
AN
          Yeast cells engineered to produce pheromone system protein surrogates,
TI
          and uses therefor
          Fowlkes, Dana M., Chapel Hill, NC, United States
IN
          Broach, Jim, Princeton, NJ, United States
          Manfredi, John, Ossining, NY, United States
          Klein, Christine, Ossining, NY, United States
Murphy, Andrew J., Montclair, NJ, United States
Paul, Jeremy, South Nyack, NY, United States
Trueheart, Joshua, South Nyack, NY, United States
Cadus Phical Corporation, Tarrytown, NY, United States (U.S.
PA
          corporation)
US 5789184
                                            19980804
                                                                                                <--
ΡI
          US 1995-464531 19950605 (8)
Continuation-in-part of Ser. No. US 1994-322137, filed on 13 Oct 1994 which is a continuation-in-part of Ser. No. US 1994-309313, filed on 20
ΑI
RLI
          Sep 1994, now abandoned which is a continuation-in-part of Ser. No. US
          1994-190328, filed on 31 Jan 1994, now abandoned which is a continuation-in-part of Ser. No. US 1993-41431, filed on 31 Mar 1993,
          now abandoned
DT
          Utility
          Granted
FS
LN.CNT
          6731
          INCLM: 435/007.310
INCL
          INCLS: 435/254.110; 435/254.200; 435/254.210
                    435/007.310
NCL
                    435/254.110; 435/254.200; 435/254.210; 435/DIG.007; 435/DIG.027
          NCLS:
IC
          [6]
          ICM: G01N033-53
          435/4; 435/7.1; 435/64; 435/252.3; 435/320.1; 435/254.21; 435/254.2;
EXF
          435/254.11
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                                USPATFULL on STN
L11
       ANSWER 94 OF 125
          1998:85817 USPATFULL
AN
          Cathepsin and methods and compositions for inhibition thereof Tung, Jay S., 2224 Semeria Ave., Belmont, CA, United States
IN
          Sinha, Sukanto, 808 Junipero Serra Blvd., San Francisco, CA, United
                     94127
          States
          McConlogue, Lisa, 283 Juanita Way, San Francisco, CA, United States
          94127
          Tatsuno, Gwen, 5910 Pinewood Rd., Oakland, CA, United States 94611
Anderson, John, 21 Bucareli Dr., San Francisco, CA, United States 941
Chrysler, Susanna, 448-1/2 San Bruno Ave., Brisbane, CA, United States
           94005
 PΙ
          US 5783434
                                            19980721
                                            19950606 (8)
          US 1995-467607
 AΙ
DT
          Utility
 FS
           Granted
 LN.CNT
          2314
           INCLM: 435/219.000
 INCL
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NCLM:
                  435/219.000
NCL
                  435/006.000; 435/212.000; 530/350.000; 536/023.100; 536/024.300
         NCLS:
IC
         [6]
         ICM: C12N009-00
         ICS: C07H021-02; C07H021-04; C12Q001-68
         530/350; 435/183; 536/23.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 95 OF 125 USPATFULL on STN
L11
                        USPATFULL
         1998:79323
AN
         cDNAs associated with ataxia-telangiectasia
TI
         Shiloh, Yosef, Tel Aviv, Israel
Tagle, Danilo A., Gaitherburg, MD, United States
Collins, Francis S., Rockville, MD, United States
RAMOT-University Authority for Applied Research & Industrial Development
Ltd., Tel Aviv, Israel (non-U.S., corporation)
IN
PA
         US 5777093
                                         19980707
PΙ
         US 1995-508836 19950728 (8)
Continuation-in-part of Ser. No. US 1995-493092, filed on 21 Jun 1995
which is a continuation-in-part of Ser. No. US 1995-441822, filed on 16
AΙ
RLI
         May 1995
DT
         Utility
FS
         Granted
LN.CNT
         1825
INCL
         INCLM: 536/023.500
                  536/023.100; 536/023.400; 435/069.100; 435/320.100; 435/325.000;
         INCLS:
                   435/252.300; 530/350.000
NCL
         NCLM:
                  536/023.500
                   435/069.100; 435/252.300; 435/320.100; 435/325.000; 530/350.000;
         NCLS:
                   536/023.100; 536/023.400
IC
          [6]
         ICM: C12N015-00
         536/23.5; 536/23.1; 536/24.1; 530/350; 514/12; 514/44; 435/320.1; 435/240.2; 435/252.3; 435/252.33; 435/69.1; 435/325; 424/93.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 96 OF 125 USPATFULL on STN
L11
         1998:57716 USPATFULL
ΑÑ
         Aptamers specific for biomolecules and methods of making
TI
         Griffin, Linda, Atherton, CA, United States
Albrecht, Glenn, Redwood City, CA, United States
Latham, John, Palo Alto, CA, United States
IN
         Latham, John, Palo Alto, CA, United States
Leung, Lawrence, Hillsborough, CA, United States
         Vermaas, Eric, Oakland, CA, United States
Toole, John J., Burlingame, CA, United States
Gilead Sciences, Inc., Foster City, CA, United States (U.S. corporation)
PA
                                          19980526
PΙ
         US 5756291
                                          19950607
                                                      (8)
AΙ
         US 1995-484192
          Continuation of Ser. No. US 1992-934387, filed on 21 Aug 1992, now
RLI
          abandoned
DT
         Utility
FS
          Granted
LN.CNT
         8242
          INCLM: 435/006.000
INCL
          INCLS: 536/023.100; 530/413.000; 935/077.000; 935/078.000
NCL
          NCLM:
                   435/006.000
          NCLS:
                   530/413.000; 536/023.100
          [6]
IC
          ICM: C12Q001-68
          ICS: C07K001-14; C07H021-04; C07H021-02
EXF 435/6; 935/77; 935/78; 530/413; 536/23.1 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 97 OF 125
                              USPATFULL on STN
L11
          1998:51432
                        USPATFULL
AN
                                         ***amyloids***
                                                             or their derivatives and use
TI
          Antibodies to .beta.-
          thereof
          Suzuki, Nobuhiro, Ibaraki, Japan
IN
          Odaka, Asano, Ibaraki, Japan
          Kitada, Chieko, Osaka, Japan
          Takeda Chemical Industries Ltd., Osaka, Japan (non-U.S. corporation)
PA
                                          19980512
PI
          US 5750349
          WO 9417197
                          19940804
          US 1994-302808
                                          19940915 (8)
ΑI
                                          19940124
          WO 1994-JP89
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19940915 PCT 102(e) date
PRAI
        JP 1993-10132
                                 19930125
                                 19930205
        JP 1993-19035
                                 19931116
        JP 1993-286985
        JP 1993-334773
                                 19931228
DT
        Utility
FS
        Granted
LN.CNT
        2609
        INCLM: 435/007.100
INCL
        INCLS: 435/007.920; 435/007.940; 435/007.950; 435/070.210; 435/326.000; 435/331.000; 530/387.900; 530/388.100; 530/389.100
                 435/007.100
        NCLM:
NCL
                 435/007.920; 435/007.940; 435/007.950; 435/070.210; 435/326.000;
        NCLS:
                 435/331.000; 530/387.900; 530/388.100; 530/389.100
IC
         [6]
        ICM: G01N033-53
        435/7.1; 435/7.92; 435/7.94; 435/70.21; 435/240.27; 435/240.26;
EXF
        435/7.95; 435/331; 435/326; 436/811; 530/387.9; 530/388.1; 530/389.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                           USPATFULL on STN
      ANSWER 98 OF 125
Lll
        1998:48435 USPATFULL
Benzylidene rhodanines
AN
TI
        Panetta, Jill A., Zionsville, IN, United States
IN
        Phillips, Michael L., Indianapolis, IN, United States
Reel, Jon K., Carmel, IN, United States
        Shadle, John K., Fishers, IN, United States
Sigmund, Sandra K., Indianpolis, IN, United States
Simon, Richard L., Greenwood, IN, United States
        Whitesitt, Celia A., Greenwood, IN, United States
        Eli Lilly and Company, Indianapolis, IN, United States (U.S.
PA
        corporation)
US 5747517
US 1996-710102
                                                                                    <--
                                      19980505
PΙ
                                      19960911 (8)
AΙ
        Division of Ser. No. US 1994-213873, filed on 16 Mar 1994
RLI
        Utility
DT
        Granted
FS
LN.CNT 2617
         INCLM: 514/369.000
INCL
         INCLS: 548/183.000
NCL
        NCLM:
                 514/369.000
        NCLS:
                 548/183.000
IC
         [6]
         ICM: C07D277-34
         ICS: A61K031-425
         548/183; 514/369
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 99 OF 125
                           USPATFULL on STN
L11
         1998:48195 USPATFULL
AN
        Method and device for diagnosing and distinguishing chest pain in early
ΤI
         onset thereof
         Jackowski, George, Inglewood, Canada
IN
         Spectral Diagnostics Inc., Toronto, Canada (non-U.S. corporation)
PA
                                       19980505
         US 5747274
PΙ
         US 1996-697690
                                      19960905 (8)
ΑI
         Continuation of Ser. No. US 1995-420298, filed on 11 Apr 1995, now
RLI
         patented, Pat. No. US 5604105 which is a continuation-in-part of Ser.
         No. US 1993-26453, filed on 3 Mar 1993, now abandoned which is a
         continuation-in-part of Ser. No. US 1991-695381, filed on 3 May 1991,
         now patented, Pat. No. US 5290678, issued on 1 Mar 1994
                                 19901012
PRAI
         CA 1990-2027434
         Utility
DT
FS
         Granted
LN.CNT
         2438
         INCLM: 435/007.940
INCL
         INCLS: 422/056.000; 422/058.000; 422/060.000; 422/061.000; 435/007.930;
                  435/007.940; 435/970.000; 435/973.000; 435/975.000; 436/514.000;
                 436/528.000; 436/530.000; 436/531.000; 436/161.000; 436/164.000; 436/807.000; 436/808.000; 436/810.000; 436/811.000
NCL
         NCLM:
                  435/007.940
                 422/056.000; 422/058.000; 422/060.000; 422/061.000; 435/007.930; 435/970.000; 435/973.000; 435/975.000; 436/161.000; 436/164.000; 436/514.000; 436/528.000; 436/530.000; 436/531.000; 436/807.000; 436/808.000; 436/810.000; 436/811.000
         NCLS:
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ICM: G01N033-573
         ICS: G01N033-558
         422/55; 422/56; 422/58; 422/60; 422/61; 435/7.9; 435/7.92; 435/7.93;
EXF
         435/7.94; 435/7.4; 435/969; 435/970; 435/973; 435/975; 436/514; 436/528; 436/530; 436/531; 436/161; 436/164; 436/807; 436/808; 436/810; 436/811
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 100 OF 125 USPATFULL on STN
L11
                         USPATFULL
AN
         1998:45097
         Method and device for diagnosing and distinguishing chest pain in early
TI
         onset thereof
IN
         Jackowski, George, Inglewood, Canada
         Spectral Diagnostics Inc., Toronto, Canada (non-U.S. corporation)
PA
                                          19980428
         US 5744358
ΡI
         US 1996-707594
                                          19960905 (8)
AΙ
         Continuation of Ser. No. US 1995-420298, filed on 11 Apr 1995, now
RLI
         patented, Pat. No. US 5604105 which is a continuation-in-part of Ser.
         No. US 1993-26453, filed on 3 Mar 1993, now abandoned which is a
         continuation-in-part of Ser. No. US 1991-695381, filed on 3 May 1991, now patented, Pat. No. US 5290678, issued on 1 Mar 1994
                                    19901012
PRAI
         CA 1990-2027434
DT
         Utility
FS
         Granted
LN.CNT
         2396
INCL
         INCLM: 435/007.400
         INCLS: 422/056.000; 422/058.000; 422/060.000; 422/061.000; 435/007.940;
                   435/970.000; 435/973.000; 435/975.000; 436/514.000; 436/528.000;
                   436/530.000; 436/531.000; 436/161.000; 436/164.000; 436/807.000; 436/808.000; 436/810.000; 436/811.000
NCL
         NCLM:
                   435/007.400
                   422/056.000; 422/058.000; 422/060.000; 422/061.000; 435/007.940; 435/970.000; 435/973.000; 435/975.000; 436/161.000; 436/164.000; 436/514.000; 436/528.000; 436/530.000; 436/531.000; 436/807.000; 436/808.000; 436/810.000; 436/811.000
         NCLS:
IC
          [6]
         ICM: G01N033-573
         ICS: G01N033-558
         422/55; 422/56; 422/58; 422/60; 422/61; 435/7.9; 435/7.92; 435/7.94; 435/7.4; 435/969; 435/970; 435/973; 435/975; 436/514; 436/528; 436/530; 436/531; 436/161; 436/164; 436/807; 436/808; 436/810; 436/811
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 101 OF 125 USPATION 1998:44877 USPATFULL
                               USPATFULL on STN
L11
AN
         Sequence-directed DNA-binding molecules compositions and methods
TI
         Edwards, Cynthia A., Menlo Park, CA, United States
IN
         Fry, Kirk E., Palo Alto, CA, United States
Cantor, Charles R., Boston, MA, United States
Andrews, Beth M., Maynard, MA, United States
PA
         Genelabs Technologies, Inc., Redwood City, CA, United States (U.S.
         corporation)
US 5744131
US 1995-476876
PΙ
                                          19980428
ΑI
                                          19950607 (8)
         Division of Ser. No. US 1992-996783, filed on 23 Dec 1992 which is a continuation-in-part of Ser. No. US 1991-723618, filed on 27 Jun 1991,
RLI
         now abandoned
DT
         Utility
FS
         Granted
LN.CNT
         5113
INCL
          INCLM: 424/078.080
          INCLS: 436/501.000; 514/001.000
                   424/078.080
NCL
         NCLM:
         NCLS:
                   436/501.000; 514/001.000
IC
          [6]
          ICM: A61K031-74
          ICS: G01N033-566; G01N033-558
EXF
          536/23.1; 536/27.1; 546/109; 436/501; 514/1; 424/78.08
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 102 OF 125 USPATFULL on STN
L11
          1998:39383 USPATFULL
AN
TI
          Sequence-directed DNA-binding molecules compositions and methods
         Edwards, Cynthia A., Menlo Park, CA, United States Fry, Kirk E., Palo Alto, CA, United States
IN
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Cantor, Charles R., Boston, MA, United States

```
Genelabs Technologies, Inc., Redwood City, CA, United States (U.S.
PA
        corporation)
                                      19980414
PΙ
        US 5738990
        US 1995-475221
                                      19950607 (8)
AΙ
        Division of Ser. No. US 1992-996783, filed on 23 Dec 1992 which is a continuation-in-part of Ser. No. US 1991-723618, filed on 27 Jun 1991,
RLI
        now abandoned
DT
        Utility
FS
        Granted
LN.CNT
        5040
        INCLM: 435/006.000
INCL
        INCLS: 435/691.000; 435/172.300; 435/320.100; 536/024.100; 935/036.000; 935/039.000
                 435/006.000
NCL
        NCLM:
                 435/069.100; 435/320.100; 536/024.100
        NCLS:
IC
        [6]
        ICM: C12P021-02
        ICS: C12N015-67; C07H021-04
        435/172.1; 435/69.1; 435/6; 435/320.1; 435/172.3; 536/24.1; 935/36;
EXF
        935/39
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 103 OF 125 USPATFULL on STN
L11
                      USPATFULL
        1998:28186
ΑN
        Mutated proteins associated with ataxia-telangiectasia Shiloh, Yosef, Tel Aviv, Israel Tagle, Danilo A., Gaitherburg, MD, United States
ΤI
IN
        Collins, Francis S., Rockville, MD, United States
Ramot-University Authority For Applied Research and Industrial
PA
        Development, Ltd., Tel Aviv, Israel (non-U.S. corporation)
PΙ
        US 5728807
                                      19980317
        US 1995-493092
                                      19950621 (8)
AΙ
        Continuation-in-part of Ser. No. US 1995-441822, filed on 16 May 1995
RLI
DT
        Utility
        Granted
FS
LN.CNT
        1637
        INCLM: 530/350.000
INCL
        INCLS:
                530/324.000; 530/326.000; 536/023.100; 536/023.500; 536/023.200
                 530/350.000
NCL
        NCLM:
                 530/324.000; 530/326.000; 536/023.100; 536/023.200; 536/023.500
        NCLS:
IC
         [6]
        ICM: C07K014-00
        ICS: C07K014-435
EXF 530/350; 530/324; 530/326; 536/23.1; 536/23.5 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 104 OF 125 USPATFULL on STN
AN
        1998:25075 USPATFULL
        Screening assay for the detection of DNA-binding molecules
TI
        Edwards, Cynthia A., Menlo Park, CA, United States Cantor, Charles R., Boston, MA, United States
IN
        Andrews, Beth M., Watertown, MA, United States
Turin, Lisa M., Berkeley, CA, United States
Genelabs Technologies, Inc., Redwood City, CA, United States (U.S.
PA
        corporation)
        US 5726014
PΙ
                                      19980310
                                                                                    <--
        US 1993-123936
                                      19930917 (8)
AΙ
        Continuation-in-part of Ser. No. US 1992-996783, filed on 23 Dec 1992
RLI
        which is a continuation-in-part of Ser. No. US 1991-723618, filed on 27
        Jun 1991, now abandoned
DT
        Utility
FS
        Granted
LN.CNT
        5659
         INCLM: 435/006.000
INCL
         INCLS: 435/091.200; 436/501.000
        NCLM:
                 435/006.000
NCL
        NCLS:
                 435/091.200; 436/501.000
         [6]
IC
         ICM: C12Q001-68
         ICS: C12P019-34; G01N033-566
         435/6; 435/235; 435/91.1; 435/91.2; 435/91.5; 536/23.1; 536/23.2;
EXF
         436/501
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 105 OF 125 USPATFULL on STN
```

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Non-crosslinked protein particles for therapeutic and diagnostic use
IN
         Yen, Richard C. K., Yorba Linda, CA, United States
         Hemosphere, Inc., Irvine, CA, United States (U.S. corporation)
PA
                                       19980310
         US 5725804
ΡI
ΑI
         US 1995-471650
                                       19950606 (8)
         Continuation-in-part of Ser. No. US 1994-212546, filed on 14 Mar 1994,
RLI
         now patented, Pat. No. US 5616311 which is a continuation-in-part of
         Ser. No. US 1993-69831, filed on 1 Jun 1993, now abandoned And Ser. No.
         US 1992-959560, filed on 13 Oct 1992, now patented, Pat. No. US 5308620 which is a continuation-in-part of Ser. No. US 1991-641720, filed on 15
         Jan 1991, now abandoned
DT
         Utility
FS
         Granted
LN.CNT
         2178
INCL
         INCLM: 252/314.000
         INCLS: 252/311.000; 424/484.000; 424/491.000; 514/776.000; 514/937.000;
                  514/965.000
NCL
         NCLM:
                  516/077.000
                  424/484.000; 424/491.000; 514/776.000; 514/937.000; 514/965.000;
         NCLS:
                  516/917.000; 516/922.000
IC
         [6]
         ICM: A61K009-64
         ICS: A61K047-42; B01J013-00
         264/4.3; 427/213.3; 427/213.33; 427/2.14; 427/2.21; 514/965; 514/937; 514/776; 252/311; 252/314; 424/491
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 106 OF 125 USPATFULL on STN
L11
         1998:14822 USPATFULL
AN
         Compounds useful as hypoglycemic agents and for treating Alzheimer's
TI
         disease
        Bue-Valleskey, Juliana M., Indianapolis, IN, United States
Hunden, David C., Carmel, IN, United States
Jones, Charles D., Indianapolis, IN, United States
Panetta, Jill A., Zionsville, IN, United States
Shaw, Walter N., Indianapolis, IN, United States
IN
         Eli Lilly and Company, Indianapolis, IN, United States (U.S.
PA
         corporation)
                                       19980210
PΙ
         US 5716975
                                                                                       <--
         US 1995-470822
                                       19950606 (8)
ΑI
         Division of Ser. No. US 1994-213651, filed on 16 Mar 1994, now patented, Pat. No. US 5523314 which is a continuation-in-part of Ser. No. US
RLI
         1992-943353, filed on 10 Sep 1992, now abandoned
DT
         Utility
FS
         Granted
LN.CNT
         1941
INCL
         INCLM: 514/369.000
         INCLS: 548/183.000
NCL
         NCLM:
                  514/369.000
         NCLS:
                  548/183.000
IC
         [6]
         ICM: C07D277-31
         ICS: A61K031-125
548/183; 514/369
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 107 OF 125 USPATFULL on STN
                       USPATFULL
AN
         1998:14634
TI
         Method of constructing sequence-specific DNA-binding molecules
         Edwards, Cynthia A., Menlo Park, CA, United States Fry, Kirk E., Palo Alto, CA, United States
IN
         Cantor, Charles R., Boston, MA, United States
Andrews, Beth M., Watertown, MA, United States
Genelabs Technologies, Inc., Redwood City, CA, United States (U.S.
PA
         corporation)
         US 5716780
PΙ
                                        19980210
                                                                                       <--
AΙ
         US 1995-484499
                                        19950607 (8)
         Division of Ser. No. US 1992-996783, filed on 23 Dec 1992 which is a
RLI
         continuation-in-part of Ser. No. US 1991-723618, filed on 27 Jun 1991,
         now abandoned
DT
         Utility
FS
         Granted
LN.CNT
        4929
INCL
         INCLM: 435/006.000
         INCLS: 436/501.000
```

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NCLS: 436/501.000
IC
          [6]
          ICM: C12Q001-68
          ICS: G01N033-566
EXF 435/6; 536/24.5; 935/33; 935/34; 935/36; 436/501 CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 108 OF 125 USPATFULL on STN
L11
          1998:6930 USPATFULL
AN
          Method and device for diagnosing and distinguishing chest pain in early
TI
          onset thereof
          Jackowski, George, Inglewood, Canada
IN
          Spectral Diagnostics Inc., Toronto, Canada (non-U.S. corporation)
PA
                                             19980120
ΡI
          UŠ 5710008
          US 1996-735178 19961022 (8)
Continuation-in-part of Ser. No. US 1995-420298, filed on 11 Apr 1995,
AΙ
RLI
          now patented, Pat. No. US 5604105 which is a continuation-in-part of Ser. No. US 1993-26453, filed on 3 Mar 1993, now abandoned which is a continuation-in-part of Ser. No. US 1991-695381, filed on 3 May 1991, now patented, Pat. No. US 5290678, issued on 1 Mar 1994
          CA 1990-2027434
                                       19901012
PRAI
DT
          Utility
          Granted
FS
LN.CNT
          2559
          INCLM: 435/007.400
INCL
          INCLS: 422/056.000; 422/058.000; 435/007.940; 435/970.000; 435/973.000; 435/975.000; 436/514.000; 436/528.000; 436/530.000; 436/807.000; 436/808.000; 436/810.000
          NCLM:
                    435/007.400
NCL
                    422/056.000; 422/058.000; 435/007.940; 435/970.000; 435/973.000; 435/975.000; 436/514.000; 436/528.000; 436/530.000; 436/807.000; 436/808.000; 436/810.000
          NCLS:
IC
          [6]
          ICM: G01N033-573
          435/7.4; 435/7.94; 435/13; 435/969; 435/970; 435/973; 435/975; 435/7.9; 435/7.92; 436/514; 436/528; 436/530; 436/541; 436/807; 436/808; 436/810;
EXF
          436/811; 422/55; 422/56; 422/58; 422/60; 422/61
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
       ANSWER 109 OF 125 USPATFULL on STN
          97:112300 USPATFULL
ΑN
          Method of ordering sequence binding preferences of a DNA-binding
TI
          molecule
          Edwards, Cynthia A., Menlo Park, CA, United States
IN
          Fry, Kirk E., Palo Alto, CA, United States
Cantor, Charles R., Boston, MA, United States
Andrews, Beth M., Maynard, MA, United States4)
          Genelabs Technologies, Inc., Redwood City, CA, United States (U.S.
PA
          corporation)
US 5693463
US 1992-996783
ΡI
                                             19971202
                                                                                                   <--
                                             19921223 (7)
ΑI
          Continuation-in-part of Ser. No. US 1991-723618, filed on 27 Jun 1991,
RLI
          now abandoned Utility
DT
FS
          Granted
LN.CNT 4908
          INCLM: 435/006.000
INCL
           INCLS: 435/007.230; 536/023.100; 935/076.000; 935/077.000
                    435/006.000
NCL
          NCLM:
          NCLS:
                    435/007.230; 536/023.100
IC
           [6]
           ICM: C12Q001-68
ICS: G01N033-574; C07H021-02; C12N015-00
EXF 435/6; 435/235; 536/23.1; 536/23.2; 514/44; 530/350; 530/351
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                                  USPATFULL on STN
       ANSWER 110 OF 125
L11
           97:106940 USPATFULL
AN
TI
           Prevention of protein aggregation
          Solomon, Beka, Herzlya, Israel
RAMOT University Authority For Applied Research and Development Ltd.,
Tel Aviv, Israel (non-U.S. corporation)
IN
PA
          US 5688651
                                              19971118
PΙ
                                                                                                    <--
                                              19941216 (8)
          US 1994-358786
AΙ
DT
          Utility
```

```
LN.CNT 1212
        INCLM: 435/007.100
INCL
        INCLS: 424/130.100; 436/063.000; 530/388.100
        NCLM:
                 435/007.100
NCL
                 424/130.100; 436/063.000; 530/388.100
        NCLS:
IC
         [6]
         ICM: G01N033-53
        ICS: G01N033-48; A61K039-395; C07K016-00 424/130.1; 424/135.1; 424/141.1; 435/7.1; 436/63; 514/44; 530/387.1; 530/388.1; 530/388.2; 530/389.1; 530/390.5
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 111 OF 125
                             USPATFULL on STN
                     USPATFULL
AN
         97:61794
        Cloning and expression of neurocan, a chondroitin sulfate proteoglycan
ΤI
IN
        Margolis, Richard U., New York, NY, United States
        Rauch, Uwe, New York, NY, United States
        Margolis, Renee K., New York, NY, United States
New York University, New York, NY, United States (U.S. corporation)
The Research Foundation of State University of New York, Albany, NY,
PA
                                                   a part interest
        United States (U.S. corporation)
        US 5648465
US 1994-340428
                                       19970715
PΙ
                                                  (8)
AΙ
                                      19941114
         Continuation of Ser. No. US 1992-922911, filed on 3 Aug 1992, now
RLI
        abandoned
DT
        Utility
        Granted
FS
LN.CNT
        2928
INCL
         INCLM: 530/350.000
         INCLS: 530/395.000; 435/069.100
NCL
        NCLM:
                 530/350.000
                 435/069.100; 530/395.000
        NCLS:
IC
         [6]
         ICM: C07K014-47
         ICS: C12N015-12
         530/350; 530/395; 514/8; 435/69.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 112 OF 125
                             USPATFULL on STN
L11
                     USPATFULL
         97:26904
AN
        Non-crosslinked protein particles for therapeutic and diagnostic use Yen, Richard C. K., Glendora, CA, United States Hemosphere, Inc., Irvine, CA, United States (U.S. corporation)
ΤI
IN
PA
                                       19970401
PΙ
         US 5616311
         US 1994-212546
                                       19940314 (8)
ΑI
         Continuation-in-part of Ser. No. US 1993-69831, filed on 1 Jun 1993, now
RLI
         abandoned And Ser. No. US 1992-959560, filed on 13 Oct 1992, now
         patented, Pat. No. US 5308620 which is a continuation-in-part of Ser.
         No. US 1991-641720, filed on 15 Jan 1991, now abandoned
DT
         Utility
FS
         Granted
LN.CNT
        2585
INCL
         INCLM: 424/001.330
         INCLS: 424/001.290; 424/001.370; 424/484.000; 424/499.000; 424/002.140;
                  424/002.210; 424/213.300; 424/213.330; 428/402.200; 428/402.240;
                  435/177.000; 935/054.000
         NCLM:
NCL
                  424/001.330
                  424/001.290; 424/001.370; 424/484.000; 424/499.000; 427/002.140;
         NCLS:
                  427/002.210; 427/213.300; 427/213.330; 428/402.200; 428/402.240;
                  435/177.000
IC
         [6]
         ICM: A61K051-08
         ICS: A61K009-50; B01J013-08; C12N011-02
264/4.3; 427/213.33; 427/2; 427/2.14; 427/2.21; 427/3; 427/213.3;
428/402.2; 428/402.24; 424/1.29; 424/1.33; 424/1.37; 424/484; 424/499;
514/832; 514/965; 935/54; 435/177
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 113 OF 125
                             USPATFULL on STN
L11
AN
         97:14582
                     USPATFULL
         Method and device for diagnosing and distinguishing chest pain in early
TI
         onset thereof
IN
         Jackowski, George, Inglewood, Canada
         Spectral Diagnostics Inc., Toronto, Canada (non-U.S. corporation)
PA
                                       19970218
         US 5604105
PΙ
```

```
Continuation-in-part of Ser. No. US 1993-26453, filed on 3 Mar 1993, now
RLI
         abandoned which is a continuation-in-part of Ser. No. US 1991-695381,
         filed on 3 May 1991, now patented, Pat. No. US 5290678, issued on 1 Mar
         1994
         CA 1990-2027434
                                     19901012
PRAI
DT
         Utility
FS
         Granted
LN.CNT
         2462
INCL
         INCLM: 435/007.400
         INCLS: 422/056.000; 422/058.000; 435/007.940; 435/970.000; 435/973.000; 435/975.000; 436/514.000; 436/528.000; 436/530.000; 436/807.000;
                   436/808.000; 436/810.000
                   435/007.400
NCL
         NCLM:
                   422/056.000; 422/058.000; 435/007.940; 435/970.000; 435/973.000;
         NCLS:
                   435/975.000; 436/514.000; 436/528.000; 436/530.000; 436/807.000;
                   436/808.000; 436/810.000
IC
          [6]
          ICM: G01N033-573
         ICS: G01N033-558
EXF 435/7.4; 435/7.9; 435/7.92; 435/7.94; 435/13; 435/969; 435/970; 435/973; 435/975; 436/528; 436/530; 436/541; 436/808; 436/810; 436/811; 422/55; 422/56; 422/58; 422/60; 422/61
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 114 OF 125 USPATFULL on STN
L11
         97:1322 USPATFULL
AN
         Dioxetane compounds for the chemiluminescent detection of proteases,
TI
         methods of use and kits therefore
         Bronstein, Irena, Newton, MA, United States
Edwards, Brooks, Cambridge, MA, United States
IN
         Martin, Christopher, Belmont, MA, United States
         Sparks, Alison, North Andover, MA, United States
Voyta, John C., Sudbury, MA, United States
Tropix, Inc., New Bedford, MA, United States (U.S. corporation)
US 5591591 19970107 <--
PA
ΡI
         US 1995-385788
                                          19950209 (8)
ΑI
         Utility
DT
FS
         Granted
LN.CNT
         747
          INCLM: 435/007.400
INCL
          INCLS: 435/006.000; 530/330.000; 530/331.000; 530/807.000; 548/526.000;
                   549/264.000; 549/332.000
NCL
         NCLM:
                   435/007.400
         NCLS:
                   435/006.000; 530/330.000; 530/331.000; 530/807.000; 548/526.000;
                   549/264.000; 549/332.000
IC
          [6]
          ICM: G01N033-573
          ICS: C07K005-06; C07K005-08; C07K005-10
          530/330; 530/331; 530/807; 435/6; 435/7.4; 548/526; 549/264; 549/332
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 115 OF 125 USPATFULL on STN
L11
AN
          96:108816
                        USPATFULL
TI
          Sequence-directed DNA-binding molecules compositions and methods
         Edwards, Cynthia A., Menlo Park, CA, United States
Cantor, Charles R., Boston, MA, United States
Andrews, Beth M., Maynard, MA, United States
Turin, Lisa M., Redwood City, CA, United States
Fry, Kirk E., Palo Alto, CA, United States
Genelabs Technologies, Inc., Redwood City, CA, United States
(U.S.
IN
PA
          corporation)
         US 5578444
US 1993-171389
PI
                                           19961126
                                                                                              <--
         US 1993-171389 19931220 (8)
Continuation-in-part of Ser. No. US 1993-123936, filed on 17 Sep 1993
AΙ
RLI
         which is a continuation-in-part of Ser. No. US 1992-996783, filed on 23 Dec 1992 which is a continuation-in-part of Ser. No. US 1991-723618,
          filed on 27 Jun 1991, now abandoned
DT
          Utility
FS
          Granted
LN.CNT
          5845
INCL
          INCLM: 435/006.000
          INCLS: 435/007.230; 536/023.100; 935/076.000; 935/077.000
NCL
          NCLM:
                   435/006.000
          NCLS:
                   435/007.230; 536/023.100
IC
          [6]
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ICS: C12N015-00; G01N033-574; C07H021-02
EXF
         435/6; 536/23.1; 536/23.2
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 116 OF 125 USPATFULL on STN
L11
         96:101466 USPATFULL
AN
         Directed evolution of novel binding proteins
TI
         Ladner, Robert C., Ijamsville, MD, United States
ΙN
         Guterman, Sonia K., Belmont, MA, United States
Roberts, Bruce L., Milford, MA, United States
Markland, William, Milford, MA, United States
Ley, Arthur C., Newton, MA, United States
Kent, Rachel B., Boxborough, MA, United States
Protein Engineering Corporation, Cambridge, MA, United States
Corporation
PA
         corporation)
PΙ
         US 5571698
                                          19961105
         US 1993-57667
                                         19930618 (8)
ΑI
         Continuation of Ser. No. US 1991-664989, filed on 1 Mar 1991, now
RLI
         patented, Pat. No. US 5223409 which is a continuation-in-part of Ser.
         No. US 1990-487063, filed on 2 Mar 1990, now abandoned which is a
         continuation-in-part of Ser. No. US 1988-240160, filed on 2 Sep 1988,
         now_abandoned
DT
         Utility
FS
         Granted
LN.CNT
         15323
INCL
         INCLM: 435/069.700
         INCLS: 435/006.000; 435/064.100; 435/172.300; 435/252.300; 435/320.100
NCL
                  435/069.700
         NCLM:
         NCLS:
                  435/006.000; 435/069.100; 435/252.300; 435/320.100; 435/477.000
IC
         [6]
         ICM: C12N025-62
         435/6; 435/64.1; 435/64.7; 435/172.3; 435/252.3; 435/320.1
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
                                USPATFULL on STN
      ANSWER 117 OF 125
L11
         96:92039
                      USPATFULL
AN
         Peptide linkage unit
TI
         Janda, Kim D., San Diego, CA, United States
Wirsching, Peter, Solana Beach, CA, United States
Ikeda, Shoji, San Diego, CA, United States
IN
         The Scripps Research Institute, La Jolla, CA, United States (U.S.
PA
         corporation)
US 5563121
PΙ
                                          19961008
         WO 9300228
                         19930111
                                                                                            <--
         US 1994-256236
                                          19940630 (8)
ΑI
         WO 1993-US228
                                          19930111
                                          19940630
                                                       PCT 371 date
                                                       PCT 102(e) date
                                          19940630
DT
         Utility
FS
         Granted
LN.CNT
         1691
INCL
         INCLM: 514/007.000
                  530/323.000; 530/326.000; 530/327.000; 530/328.000; 530/329.000; 530/330.000; 562/017.000; 562/018.000; 930/030.000
         INCLS:
NCL
                   514/007.000
         NCLM:
         NCLS:
                   530/323.000; 530/326.000; 530/327.000; 530/328.000; 530/329.000;
                   530/330.000; 562/017.000; 562/018.000; 930/030.000
IC
          [6]
         ICM: A61K038-03
         ICS: C07K004-00; C07K005-02; C07K007-02
514/2; 514/14; 514/15; 514/16; 514/17; 514/18; 514/7; 930/21; 930/30;
530/323; 530/326; 530/327; 530/328; 530/329; 530/330; 530/331; 530/332;
562/17; 562/18
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
       ANSWER 118 OF 125
                                USPATFULL on STN
         96:48400
                      USPATFULL
AN
         Compounds useful as hypoglycemic agents and for treating Alzheimer's
TI
         disease
         Bue-Valleskey, Juliana M., Indianapolis, IN, United States
Hunden, David C., Carmel, IN, United States
IN
         Jones, Charles D., Indianapolis, IN, United States Panetta, Jill A., Zionsville, IN, United States Shaw, Walter N., Indianapolis, IN, United States
PA
         Eli Lilly and Company, Indianapolis, IN, United States (U.S.
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US 5523314
                                   19960604
ΡI
        US 1994-213651
                                   19940316
                                             (8)
ΑI
        Continuation-in-part of Ser. No. US 1992-943353, filed on 10 Sep 1992,
RLI
        now abandoned
        Utility
DT
FS
        Granted
LN.CNT
       2068
INCL
        INCLM: 514/369.000
        INCLS: 548/183.000
NCLM: 514/369.000
NCL
        NCLM:
        NCLS:
                548/183.000
IC
        [6]
        ICM: A61K031-425
        514/369
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 119 OF 125
                           USPATFULL on STN
L11
AN
        96:38884
                   USPATFULL
TI
        Immunological activity of rhamnolipids
        Piljac, Goran, 2323 Shasta Dr., Apt 40, Davis, CA, United States
IN
                                                                                   95616
                Visnja, 2323 Shasta Dr., Āpt 40, Davis, CA, United States 661 19960507 <--
                                                                                    95616
       US 5514661
US 1995-520076
PΙ
                                   19950828 (8)
ΑI
        Division of Ser. No. US 1994-277975,
                                                filed on 20 Feb 1994, now patented,
RLI
        Pat. No. US 5466675 which is a continuation-in-part of Ser. No. US
        1992-866691, filed on 10 Apr 1992, now abandoned
DT
        Utility
FS
        Granted
LN.CNT
        1424
INCL
        INCLM: 514/025.000
        INCLS: 514/814.000; 514/861.000; 514/863.000; 514/864.000; 514/878.000;
                514/883.000; 514/885.000; 514/886.000; 514/887.000; 514/889.000;
                514/903.000; 514/908.000
                514/025.000
NCL
        NCLM:
        NCLS:
                514/814.000; 514/861.000; 514/863.000; 514/864.000; 514/878.000;
                514/883.000; 514/885.000; 514/886.000; 514/887.000; 514/889.000;
                514/903.000; 514/908.000
IC
        [6]
        ICM: A61K031-715
        514/25; 514/814; 514/861; 514/863; 514/864; 514/878; 514/883; 514/885;
EXF
        514/886; 514/887; 514/889; 514/903; 514/908
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 120 OF 125 USPATFULL on STN
L11
        96:27100
                   USPATFULL
\mathbf{A}\mathbf{N}
TI
        Production of peptide amides
IN
        Bibbs, Jeffrey A., San Diego,
                                         CA, United States
        Lehman De Gaeta, Laura S., Olivenhain, CA, United States
Jones, Howard, Poway, CA, United States
PA
        Amylin Pharmaceuticals, Inc., San Diego, CA, United States (U.S.
        corporation)
US 5503989
US 1992-927755
                                   19960402
PI
                                                                             <--
AΙ
                                   19920810 (7)
        Continuation-in-part of Ser. No. US 1991-742768, filed on 8 Aug 1991, now abandoned And a continuation-in-part of Ser. No. US 1991-742769,
RLI
        filed on 8 Aug 1991, now abandoned
DT
        Utility
        Granted
FS
LN.CNT
        712
INCL
        INCLM: 435/068.100
        INCLS: 530/307.000; 530/309.000; 530/313.000; 530/317.000; 530/324.000;
                530/345.000
                435/068.100
530/307.000; 530/309.000; 530/313.000; 530/317.000; 530/324.000;
NCL
        NCLM:
        NCLS:
                530/345.000
IC
        [6]
        ICM: C12P021-06
        ICS: C07K005-00; C07K007-00; C07K017-00
EXF
        530/324; 530/345; 530/307; 530/317; 530/313; 530/309; 435/68.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L11
      ANSWER 121 OF 125
                          USPATFULL on STN
        95:101209
                    USPATFULL
\mathbf{A}\mathbf{N}
TI
        Immunological activity of rhamnolipids
IN
        Piljac, Goran, 2323 Shasta Dr., Apt. 40, Davis, CA, United States
                                                                                    95616
```

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95616
         US 5466675
PΙ
                                        19951114
         US 1994-277975
                                        19940720
                                                    (8)
AΙ
         Continuation-in-part of Ser. No. US 1992-866691, filed on 10 Apr 1992,
RLI
         now abandoned
PRAI
         BE 1992-115
                                   19920204
DT
         Utility
         Granted
FS
LN.CNT
         1443
         INCLM: 514/025.000
INCL
                  514/814.000; 514/861.000; 514/863.000; 514/864.000; 514/878.000; 514/883.000; 514/885.000; 514/886.000; 514/887.000; 514/889.000;
         INCLS:
                  514/903.000; 514/908.000
NCL
         NCLM:
                  514/025.000
                  514/814.000; 514/861.000; 514/863.000; 514/864.000; 514/878.000;
         NCLS:
                  514/883.000; 514/885.000; 514/886.000; 514/887.000; 514/889.000;
                  514/903.000; 514/908.000
IC
         [6]
         ICM: A61K031-715
         514/25; 514/861; 514/863; 514/864; 514/878; 514/883; 514/885; 514/886;
EXF
514/887; 514/889; 514/903; 514/908; 514/814
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 122 OF 125
                              USPATFULL on STN
L11
         95:52252 USPATFULL
AN
TI
         Amyloidin protease and uses thereof
         Dovey, Harry F., Pacifica, CA, United States
IN
         Seubert, Peter A., San Mateo, CA, United States
         Sinha, Sukanto, San Francisco, CA, United States
         Athena Neurosciences, Inc., So. San Francisco, CA, United States (U.S.
PA
         corporation)
         Eli Lilly and Company, Indianapolis, IN, United States (U.S.
         corporation)
US 5424205
                                        19950613
PI
         US 1993-59032
                                        19930507 (8)
ΑI
         Division of Ser. No. US 1991-766351, filed on 30 Sep 1991, now patented,
RLI
         Pat. No. US 5292652 which is a continuation-in-part of Ser. No. US
         1990-594122, filed on 5 Oct 1990, now abandoned
DT
         Utility
FS
         Granted
LN.CNT
         1528
         INCLM: 435/226.000
INCLS: 435/219.000
NCLM: 435/226.000
NCLS: 435/219.000
INCL
NCL
         [6]
IC
         ICM: C12N009-64
         435/226; 435/219
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 123 OF 125
                              USPATFULL on STN
L11
         94:102315
                      USPATFULL
AN
TI
         Amylin peptides
        Cooper, Garth J. S., Solana Beach, CA, United States
Willis, Antony C., Witney, England
Amylin Pharmaceuticals, Inc., San Diego, CA, United States (U.S.
IN
PA
         corporation)
PΙ
                                        19941122
         US 5367052
                                                                                         <--
        US 1989-346624 19890501 (7)
Continuation-in-part of Ser. No. US 1988-275319, filed on 23 Nov 1988, now abandoned And a continuation-in-part of Ser. No. US 1988-236985, filed on 26 Aug 1988, now abandoned, said Ser. No. US -275319 which is a continuation-in-part of Ser. No. US 1988-186520, filed on 27 Apr
ΑI
RLI
                                                                                  -275319 which
         1988, now abandoned
         GB 1987-9871
PRAI
                                   19870427
         GB 1987-20115
                                   19870826
         Utility
DT
FS
         Granted
LN.CNT
         777
         INCLM: 530/307.000
INCL
         INCLS: 530/324.000; 530/387.900
NCL
         NCLM:
                  530/307.000
         NCLS:
                  530/324.000; 530/387.900
         [5]
IC
         ICM: A61K037-02
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424/85.8; 530/307; 530/324; 514/12
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 124 OF 125 USPATFULL on STN
L11
          94:20087 USPATFULL
AN
TI
          Amyloidin protease and uses thereof
          Dovey, Harry F., Pacifica, CA, United States
Seubert, Peter A., San Mateo, CA, United States
Sinha, Sukanto, San Francisco, CA, United States
Athena Neurosciences, Inc., South San Francisco, CA, United States (U.S.
IN
PA
          corporation)
          Eli Lilly and Company, Indianapolis, IN, United States (U.S. corporation)
US 5292652 19940308
US 1991-766351 19910930 (7)
PΙ
                                                                                                     <--
ΑI
          Continuation-in-part of Ser. No. US 1990-594122, filed on 5 Oct 1990,
RLI
          now abandoned
DT
          Utility
FS
          Granted
LN.CNT
          1462
          INCLM: 435/226.000
INCL
          INCLS: 435/219.000
NCLM: 435/226.000
NCLS: 435/219.000
NCL
           [5]
IC
          ICM: C12N009-64
          435/219; 435/226; 435/23
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       ANSWER 125 OF 125 USPATFULL on STN
L11
          93:52487 USPATFULL
AN
          Directed evolution of novel binding proteins
Ladner, Robert C., Ijamsville, MD, United States
Guterman, Sonia K., Belmont, MA, United States
Roberts, Bruce L., Milford, MA, United States
Markland, William, Milford, MA, United States
TI
IN
          Ley, Arthur C., Newton, MA, United States
Kent, Rachel B., Boxborough, MA, United States
          Protein Engineering Corp., Cambridge, MA, United States (U.S.
PA
          corporation)
          US 5223409
US 1991-664989
PΙ
                                              19930629
                                                                                                     <--
          US 1991-664989 19910301 (7)
Continuation-in-part of Ser. No. US 1990-487063, filed on 2 Mar 1990, now abandoned And a continuation-in-part of Ser. No. US 1988-240160,
AΙ
RLI
          filed on 2 Sep 1988, now abandoned
          Utility
DT
FS
          Granted
LN.CNT
          15410
          INCLM: 435/069.700
INCL
          INCLS: 435/069.100; 435/172.300; 435/252.300; 435/320.100; 530/380.300;
                    530/387.500
NCL
          NCLM:
                    435/069.700
                    435/005.000; 435/069.100; 435/252.300; 435/320.100; 435/472.000; 530/387.300; 530/387.500
          NCLS:
IC
           [5]
          ICM: C12N015-09
          ICS: C12N015-62; C12N015-63
           435/69.1; 435/172.3; 435/252.3; 435/320.1; 530/350
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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STN INTERNATIONAL LOGOFF AT 16:05:07 ON 26 JAN 2005